administered orally.

- 3. (Original) The method of claim 2, wherein said conjugated linoleic acid is administered in a unit dosage form.
- 4. (Original) The method of claim 3, wherein said unit dosage form is a food product.
- 5. (Original) The method of claim 1, wherein said conjugated linoleic acid is selected from the group consisting of 9,11-octadecadienoic acid, esters thereof, geometric isomers thereof, salts thereof and mixtures thereof.
- 6. (Original) The method of claim 5, wherein said geometric isomers have configurations selected from the group consisting of trans, trans; cis, cis; trans, cis; and cis, trans.
- 7. (Original) The method of claim 1, wherein said conjugated linoleic acid is selected from the group consisting of 10,12-octadecadienoic acid, esters thereof, geometric isomers thereof, salts thereof and mixtures thereof.
- 8. (Original) The method of claim 7, wherein said geometric isomers have configurations selected from the group consisting of trans, trans; cis, cis; trans, cis; and cis, trans.
- 9. (Previously amended) The method of claim 1, wherein said conjugated linoleic acid is comprised predominantly of cis,trans-9,11-octadecadienoic acid and trans,cis-octadecadienoic acid.
- 10. (Previously amended) The method of claim 1, wherein said conjugated linoleic acid is comprised predominantly of cis, cis-9,11-octadecadienoic acid.

- 11. (Original) The method of claim 1, wherein said conjugated linoleic acid is administered in an amount of about 1 mg of said conjugated linoleic acid/kg body weight to about 10,000 mg of said conjugated linoleic acid/kg body weight.
 - 12. (Original) The method of claim 1, wherein said animal is a mammal.
 - 13. (Original) The method of claim 12, wherein said mammal is a human.
- 14. (Original) The method of claim 1. wherein said conjugated linoleic acid is administered in a pharmaceutically acceptable carrier medium.
- 15. (Original) The method of claim 14, wherein said pharmaceutically acceptable carrier medium includes water.
- 16. (Original) A food composition useful in treating diabetes comprising, a food product having a therapeutically effective amount of conjugated linoleic acid, said conjugated linoleic acid predominantly comprised of a mixture of cis,trans-9,11-octadecadienoic acid and trans,cis-9,11-octadecadienoic acid.
- 17. (Original) The food composition of claim 16, wherein said therapeutically effective amount of said mixture is sufficient to provide about 1 mg of said mixture/kg body weight per serving to about 10,000 mg of said mixture/kg body weight per serving.
- 18. (Original) A food composition useful in treating diabetes comprising, a food product having a therapeutically effective amount of conjugated linoleic acid, said conjugated linoleic acid predominantly comprised of cis,cis-9,11-octadecadienoic acid.

- 19. (Original) The food composition of claim 18, wherein said conjugated linoleic acid is administered in an amount sufficient to provide about 1 mg of said cis,cis-9,11-ocatdecadienoic acid/kg body weight per serving to about 10,000 mg of said cis,cis-9,11-octadecadienoic acid/kg body weight per serving.
- 20. (Original) A food composition useful in treating diabetes comprising, a food product having a therapeutically effective amount of conjugated linoleic acid, said conjugated linoleic acid predominantly comprised of trans, cis-10,12-octadecadienoic acid.
- 21. (Original) The food composition of claim 20, wherein said conjugated linoic acid is administered in an amount sufficient to provide about 1 mg of said trans, cis-10,12-octadecadienoic acid/kg body weight per serving to about 10,000 mg of said trans, cis-10,12-octadecadienoic acid / kg body weight per serving.
- 22. (Previously added) A method of treating symptoms of diabetes in a human comprising:
 - a) providing
 - i) a therapeutically effective amount of conjugated linoleic acid; and
 - ii) a human patient suffering from diabetes; and
- b) administering said therapeutically effective amount of conjugated linoleic acid to said human diabetic patient under conditions such that said symptoms of diabetes are treated.

Remarks

Claims 1-22 remain pending in the present application. In this response, no amendment to claims 1-22 is made. Applicants consider the claims allowable in their present form. The Examiner maintains his rejection that claims 1-22 are unpatentable as being obvious under 35 U.S.C. ¶103 over the cited references. For the reasons which are presented in the sections which